	Type	L #	Hits	Search Text	DBs
1	BRS	L1	13277	microfluid\$9	US- PGPUB; USPAT
2	BRS	L2	766	1 and (radial or circular or round) near8 (base or platform or substrate or housing or body)	US- PGPUB; USPAT
3	BRS	L3	442	2 and (pump or micropump)	US- PGPUB; USPAT
4	BRS	L4	500	2 and (capillary or capillary near8 tube)	US- PGPUB; USPAT
5	BRS	L5	332	3 and (capillary or capillary near8 tube)	US- PGPUB; USPAT
6	BRS	L6	403	2 and (disk or disc)	US- PGPUB; USPAT
7	BRS	L7	198	5 and 6	US- PGPUB; USPAT
8	BRS	L8	246	3 and 6	US- PGPUB; USPAT
9	BRS	L9	558	microfluid\$9	EPO
10	BRS	L10	0	9 and (radial or circular or round) near8 (base or platform or substrate or housing or body)	EPO
11	BRS	L11	1	9 and (radial or circular or round or disc or disk) near8 (base or platform or substrate or housing or body)	EPO
12	BRS	L12	1090	1 and (radial or circular or round or disc or disk) near8 (base or platform or substrate or housing or body)	US- PGPUB; USPAT
13	BRS	L13	569	12 and (pump or micropump)	US- PGPUB; USPAT

	Type	L #	Hits	Search Text	DBs
14	BRS	L14	418	13 and (capillary or capillary near8 tube)	US- PGPUB; USPAT
15	BRS	L15	683	12 and (capillary or capillary near8 tube)	US- PGPUB; USPAT
16	BRS	L16	2092	microfluid\$9	DERWEN T
17	BRS	L17	9	16 and (radial or circular or round or disc or disk) near8 (base or platform or substrate or housing or body)	DERWEN T
18	BRS	L19	o	18 and (radial or circular or round or disc or disk) near8 (base or platform or substrate or housing or body)	IBM_TD B
19	BRS	L18	3	microfluid\$9	IBM_TD B
20	BRS	L20	79	microfluid\$9	JPO
21	BRS	L21	0	20 and (radial or circular or round or disc or disk) near8 (base or platform or substrate or housing or body)	JPO
22	BRS	L22	0	20 and (radial or circular or round or disc or disk) with (base or platform or substrate or housing or body)	JPO

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                 truncation
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                 CA(SM)/CAplus(SM) display of CA Lexicon enhanced
NEWS 20
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                 CAS REGISTRY (SM) updated with amino acid codes for pyrrolysine
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         SEP 25
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         SEP 28
                 CEABA-VTB classification code fields reloaded with new
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L1 17691 MICROFLUID?

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L2 40 L1 AND (RADIAL? OR CIRCULAR OR ROUND) (S) (BASE OR PLATFORM OR SUBSTRATE OR HOUSING OR BODY)

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L3 ANSWER 1 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

2006:922134 CAPLUS

DOCUMENT NUMBER:

145:317317

TITLE:

Flow 'switching on a multi-structured

microfluidic cd (compact disc) using coriolis

force

INVENTOR(S):

Zoval, Jim V.; Madou, Marc J.; Jia, Guangyao; Kim,

Jitae; Kido, Horacio

PATENT ASSIGNEE(S):

The Regents of the University of California, USA

SOURCE: PCT Int. Appl., 22pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.				KIND		DATE		ž	APPLICATION NO.					DATE		
,																
WO 2006093978				A2 20060908		WO 2006-US7119				20060228						
W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	ΑZ,	ВA,	BB,	BG,	BR,	BW,	BY,	ΒZ,	CA,	CH,
	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,
						ID,										
						LT,										
	MZ,	NA,	NG,	NI,	NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,
	SG,	SK,	SL,	SM,	SY,	TJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,
	VN,	YU,	ZA,	ZM,	ZW											

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRIORITY APPLN. INFO.:

US 2005-657760P P 20050302

A microfluidic switching device includes a planar substrate having a central axis of rotation and a radially -oriented microchannel disposed in the planar substrate that terminates at a junction. In one aspect, the junction is formed as a double-layered junction in which an upstream portion is vertically offset from a downstream portion. The upstream portion has a smaller effective cross-sectional area than the downstream portion. First and 2nd outlet chambers are coupled at one end to the junction. The device is rotated about the central axis in a clockwise direction so as to cause the fluid in the reservoir to flow into the 1st (right) outlet chamber or in a counter-clockwise direction so as to cause the fluid in the reservoir to flow into the 2nd (left) outlet chamber.

L3 ANSWER 2 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER: 2006:9064103 INSPEC

TITLE: Quantification and prediction of jet macro-mixing

times in static microwell plates

AUTHOR: Lye, G.J.; Nealon, A.J.; (Dept. of Biochem. Eng.,

Univ. Coll. London, UK), O'Kennedy, R.D.;

Titchener-Hooker, N.J.

SOURCE: Chemical Engineering Science (Aug. 2006), vol.61,

no.15, p. 4860-70, 43 refs. CODEN: CESCAC, ISSN: 0009-2509

SICI: 0009-2509(200608)61:15L.4860:QPMM;1-F

Doc.No.: S0009-2509(06)00108-4 Published by: Elsevier, UK

DOCUMENT TYPE: Journal

TREATMENT CODE: Practical; Theoretical; Experimental

COUNTRY: United Kingdom

LANGUAGE: English

AN 2006:9064103 INSPEC

Automated experimentation in microwell plate formats is widely used in AB high throughput drug discovery. Such approaches are now being considered for the study of bioprocess unit operations in order to speed the delivery of new medicines to market. The generation of useful design data from microwell formats requires an understanding of the engineering environment within individual microwells. Rapid and efficient macro-mixing is crucial in this respect to ensure the generation of quantitative and reproducible data. In this study, we have developed a high-speed video technique for the accurate quantification of jet macro-mixing times in static microwell plates which also enables visualisation of jet formation and liquid flow patterns within wells. Mixing times have been determined using both the fixed (di=0.54 mm) and disposable (di=0.6 mm) tips of a Perkin Elmer MultiProbe IITM liquid handling robot for a range of jet Reynolds numbers (Rej=1000-3960) and liquid addition volumes (VA=10-859 µl). Three microwell geometries have been investigated; one that is identical to a single well from a standard 96-round well plate (Vi=200 µl) and two novel designs based upon theories of jet mixing (Vi=200 and 1720 µl). For conditions where macro-mixing was complete within the lifespan of the jet, t95 mixing times for the standard round well were in the range 0.033-0.121s while for the larger of the two designed wells they were in the range 0.228-0.705s. The rapid mixing times in the standard round well are a consequence of increased energy dissipation as the liquid jet impinges on the base of the well. For the two designed wells maximising the jet length to nozzle diameter ratio (X/di) is shown to promote the most efficient macro-mixing due to entrainment and circulation of the bulk liquid in the well. For low volume additions

and short jet lifespans it is also shown that mixing times can be of the order of minutes. Finally, the t95 results for each of the well geometries have been correlated to the conditions used for jet formation using a correlation of the form first proposed by Baldyga and co-workers [Baldyga, J., Bourne, J.R., Dubuis, B., Etchells, A.W., Gholap, R.V., Zimmermann, B., 1995. Jet reactor scale-up for mixing controlled reactions. Chemical Engineering Research & Design 73, 497-502]. This enables good prediction of the experimentally determined mixing times and estimation of the minimum liquid addition volume (VCrit) that will ensure rapid and efficient macro-mixing. The correlation therefore enables automation users to optimise or control macro-mixing times in microwell experiments. [All rights reserved Elsevier]

L3 ANSWER 3 OF 35 COMPENDEX COPYRIGHT 2006 EEI on STN DUPLICATE 1

ACCESSION NUMBER: 2006(31):3477 COMPENDEX

TITLE: Dual independent displacement-amplified micropumps

with a single actuator.

AUTHOR: Tracey, M.C. (Science and Technology Research

Institute University of Hertfordshire, Hatfield, Hertfordshire AL10 9AB, United Kingdom); Johnston,

I.D.; Davis, J.B.; Tan, C.K.L.

SOURCE: Journal of Micromechanics and Microengineering v 16 n

8 Aug 1 2006 2006.p 1444-1452, arn: 002

CODEN: JMMIEZ ISSN: 0960-1317 E-ISSN: 1361-6439

PUBLICATION YEAR: 2006
DOCUMENT TYPE: Journal
TREATMENT CODE: Theoretical
LANGUAGE: English

AN 2006(31):3477 COMPENDEX

AB We report a dual-micropump structure operated by a single actuator element. The constituent micropumps are a form of micro throttle pump (MTP) comprising a narrow flow channel incorporating two microthrottles. We term this a 'linear MTP' (LMTP). The LMTP's narrowness, in conjunction with an elastomeric substrate, allows multiple, independent, LMTPs to be actuated by a single piezoelectric actuator thereby suiting it to parallel microfluidic architectures. Furthermore, LMTP elements can be combined into parallel or series composites yielding increased maximum pumping rates or back pressures, respectively, when compared to a single LMTP element. The LMTP's flow-channel-like, linear pump chamber minimizes the development of recirculatory flows associated with circular pump chambers which, in part, determine their frequency response and hence maximum pumping rates. We have modelled, fabricated and evaluated a dual-LMTP. We report operation in three modes: as two distinct pumps, as a series composite pump, and as a parallel composite pump. Operating at about 1.6 kHz, with both pumps under identical load conditions, each pump yielded maximum pumping rates of about 750 nul min -1 and back pressures of 18 kPa, both with close matching. Configured as a series composite, a 35 kPa back pressure was achieved, and configured as a parallel composite, a maximum pumping rate of 1.4 ml min-1 resulted. Images of 5 num polystyrene beads flowing within an LMTP confirm minimal recirculatory behaviour consistent with the LMTP's increased operating frequencies compared to circular pump chamber MTPs. \$CPY 2006 IOP Publishing Ltd. 13 Refs.

L3 ANSWER 4 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2006:351297 CAPLUS

DOCUMENT NUMBER: 144:447289

TITLE: Circular orientation of biomolecular rails using micro

fluidic channels

AUTHOR(S): Watanabe, Yasumasa; Suzuki, Kenji; Takeuchi, Shoji CORPORATE SOURCE: Inst. Ind. Sci., The University of Tokyo, Japan

SOURCE: Seisan Kenkyu (2006), 58(2), 138-141

CODEN: SEKEAI; ISSN: 0037-105X

PUBLISHER: Tokyo Daigaku Seisan Gijutsu Kenkyusho

DOCUMENT TYPE: Journal LANGUAGE: Japanese

AB A microfluidic device with circular orientation of

microtubule-kinesin system was fabricated by using polydimethylsiloxane (PDMS) and glass plates. A clockwise circular orientation of microtubules was observed by gliding assay in the PDMS channels.

L3 ANSWER 5 OF 35 COMPENDEX COPYRIGHT 2006 EEI on STN

ACCESSION NUMBER: 2006(22):12262 COMPENDEX

TITLE: Fabrication of long microchannels with circular cross

section using astigmatically shaped femtosecond laser

pulses and chemical etching.

AUTHOR: Maselli, Valeria (Istituto di Fotonica e

Nanotecnologie-CNR ULTRAS-INFM-CNR Politecnico di Milano, 20133 Milan, Italy); Osellame, Roberto; Cerullo, Giulio; Ramponi, Roberta; Laporta, Paolo;

APPLICATION NO.

DATE

Magagnin, Luca; Cavallotti, Pietro Luigi

SOURCE: Applied Physics Letters v 88 n 19 2006., arn: 191107

CODEN: APPLAB ISSN: 0003-6951

PUBLICATION YEAR: 2006
DOCUMENT TYPE: Journal
TREATMENT CODE: Theoretical
LANGUAGE: English
AN 2006(22):12262 COMPENDEX

AB We report on the fabrication of microfluidic channels in fused silica using femtosecond laser irradiation followed by chemical etching. Using an astigmatically shaped beam, we achieve microchannels with circular cross section and length up to 1.5 mm. We use the same femtosecond laser, with different irradiation parameters, to fabricate high quality optical waveguides on the same substrate. The integration of microchannels and waveguides will enable a forthcoming class of biophotonic sensors. \$CPY 2006 American Institute of Physics. 14 Refs.

L3 ANSWER 6 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2005:1204677 CAPLUS

DOCUMENT NUMBER: 143:455495

TITLE: Spiral-type flow-through polymerase chain reaction

(PCR) chip

INVENTOR(S): Chen, Wenyuan; Jia, Xiaoyu; Niu, Zhiqiang; Zhang,

Weiping

KIND

PATENT ASSIGNEE(S): Shanghai Jiao Tong University, Peop. Rep. China

DATE

SOURCE: Faming Zhuanli Shenqing Gongkai Shuomingshu, 11 pp.

CODEN: CNXXEV

DOCUMENT TYPE: Patent LANGUAGE: Chinese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.

CN 1584043	A 2	20050223	CN 2004-10024703	20040527
PRIORITY APPLN. INFO.:			CN 2004-10024703	20040527
AB The invention discle	oses a sp	oiral-type f	low-through PCR o	chip for use in
fields of preventive	e medicir	ne and publi	c health. The cl	nip includes a
substrate bearing m	icrofluid	dic conduits	and a sealing ca	ap which
is sealed together w	with the	substrate b	y bonding technic	que, wherein the
microfluidic conduit	ts on the	substrate	are spiral-type;	a sample
inlet and a sample of	outlet ar	ce disposed	in the sealing ca	ap which resp.
correspond to the st	tarting a	and ending p	oints of the mich	rofluidic
conduits. A circula	ar heater	s for heati	ng the denaturat:	ion zone
and the elongation	zone, and	d a circular	temperature sens	sor are
disposed on the other	er side o	of the subst	rate opposite to	the
sealing cap; an inst				

the substrate; and a radiating zone formed by metallic film for cooling the renaturation zone is disposed in the low-right area of the chip, between the insulating layer and the substrate. The invention has the advantages of increased efficiency of amplification, and broadened application range, enhanced efficiency, reduced size, as well as improved compatibility and integratability of the chip.

L3 ANSWER 7 OF 35 COMPENDEX COPYRIGHT 2006 EEI on STN

ACCESSION NUMBER: 2006(17):1999 COMPENDEX

TITLE: Building a tissue engineered microfluidic

bioreactor array for high-throughput assays.

AUTHOR: Wen, Yuan; Yang, Shang-Tian

MEETING TITLE: 05AICHE: 2005 AICHE Annual Meeting and Fall Showcase.

MEETING LOCATION: Cincinnati, OH, United States

MEETING DATE: 30 Oct 2005-04 Nov 2005

SOURCE: AIChE Annual Meeting, Conference Proceedings 2005.p

12678

SOURCE: 05AIChE: 2005 AIChE Annual Meeting and Fall Showcase,

Conference Proceedings

PUBLICATION YEAR: 2005 MEETING NUMBER: 66925

DOCUMENT TYPE: Conference Article

TREATMENT CODE: Theoretical; Experimental

LANGUAGE: English
AN 2006(17):1999 COMPENDEX

As the progress of functional genomics, the number of drug candidates has AB by far outgrown the present capability of cytotoxicity studies. In addition, early discoveries of toxic side effects of drugs in test can greatly reduce the risk of a lengthy and expensive drug development process. Increasing interest in toxicity study systems in vitro has made cell-based platforms an attractive approach to uncovering the toxicity effects at cellular and subcellular levels. However, conventionally, cells cultured statically on 2D surfaces may not well exhibit authentic responses upon external stimulation. We have designed and fabricated a mcirofluidic device using multiple layers of poly(dimethylsiloxane) (PDMS) through photolithography and replica molding. Each layer was designed to specifically serve part of a network of microfluidic channels for medium flow, drug serial dilution, mixers and cell culture chambers with the same dimension as the round wells of a 384 well plate. Tissue engineering scaffolds were placed in the chambers for three dimensional cell cultures on the chip. The alignment and bonding of different PDMS layers were investigated. The fluid ports were also engineered for prolonged medium flow and for convenience of the chip to be scanned within a 384 well plate reader. Optimal designs for uniform fluid distribution in the highly parallel system were also studied. Computational fluidic dynamics was also explored for the simulation of the flow rate and mass transport properties of serial dilution channels, cell culture chambers, mixers and overall fluid distribution with FluentTM. The microfluidic bioreactor array was designed to test the effects of 6 different concentrations of a drug with controls on two different types of cells in a perfusion 3D culture without interference. The numbers of cell types and drugs for the test can be easily expanded with similar designs. In this study, embryonic stem cells stably transfected with fluorescent proteins cultured in the microfluidic bioreactor array were studied for high throughput assays of drugs. The test results based on the cellular fluorescence intensity change caused by the drugs will be presented in the paper.

L3 ANSWER 8 OF 35 COMPENDEX COPYRIGHT 2006 EEI on STN

ACCESSION NUMBER: 2005(49):5575 COMPENDEX

TITLE: Thermoplastic microfluidic platform for

single-molecule detection, cell culture, and

actuation.

AUTHOR: Melin, Jenas (Beijer Laboratory Department of Genetics

and Pathology Hudbeck Laboratory, Uppsala, Sweden);

Johansson, Henrik; Soderberg, Ola; Nikolajeff,

Fredrik; Landegren, Ulf; Nilsson, Mats; Jarvius, Jonas

SOURCE: Analytical Chemistry v 77 n 22 Nov 15 2005 2005.p

7122-7130

CODEN: ANCHAM ISSN: 0003-2700

PUBLICATION YEAR: 2005
DOCUMENT TYPE: Journal
TREATMENT CODE: Theoretical
LANGUAGE: English
AN 2005(49):5575 COMPENDEX

We have developed a multipurpose microfluidic platform AB that allows for sensitive fluorescence detection on inexpensive disposable chips. The fabrication scheme involves rapid injection molding of thermoplastics, followed by silica deposition and covalent attachment of an unstructured flexible lid. This combines the virtues of elastomer technology with high-throughput compact disk injection molding. Using this technique, the time to produce 100 chips using a single master can be lowered from more than 1 week by standard PDMS technologies to only a couple hours. The optical properties of the fabricated chips were evaluated by studying individual fluorescence-labeled DNA molecules in a microchannel. Concatemeric DNA molecules were generated through rolling circle replication of circular DNA molecules, which were labeled by hybridization of fluorescence-tagged oligonucleotides. Rolling circle products (RCPs) were detected after as little as 5 min of DNA polymerization, and the RCPs in solution showed no tendency for aggregation. To illustrate the versatility of the platform, we demonstrate two additional applications: The flexible property of the lid was used to create a peristaltic pump generating a flow rate of 9 nL/s. Biocompatibility of the platform was illustrated by culturing Chinese hamster ovary cells for 7 days in the microfluidic

L3 ANSWER 9 OF 35 COMPENDEX COPYRIGHT 2006 EEI on STN

channels. \$CPY 2005 American Chemical Society. 29 Refs.

ACCESSION NUMBER:

2004(52):8757 COMPENDEX

TITLE:

Hybridization enhancement using microfluidic

planetary centrifugal mixing.

AUTHOR:

Bynum, Magdalena A. (Agilent Laboratories Agilent Technologies, Palo Alto, CA 94304, United States);

Gordon, Gary B.

SOURCE:

Analytical Chemistry v 76 n 23 Jan 3 2005 2005.p

7039-7044

CODEN: ANCHAM ISSN: 0003-2700

PUBLICATION YEAR: 2005
DOCUMENT TYPE: Journal
TREATMENT CODE: Experimental
LANGUAGE: English

2004(52):8757 COMPENDEX AN DNA microarrays produce their greatest sensitivities when hybridized using AB concentrated samples and effective mixing; however, these goals have proved elusive to combine. If samples are diluted enough to fill larger chambers, then mixing works well using either pumping or gravity with rotation, although sensitivities will suffer. Various techniques for mixing concentrated samples in small thin chambers have been proposed; however, they often leave streaks or scars, and their reusable components require careful cleaning. Here we introduce a versatile new microfluidics platform, a two-axis centrifuge whose fluidic chambers rotate in a planetary relationship to a radial gravitational field. This paradigm readily overcomes surface and viscous forces even in chambers only 50 mum thin. Thin chambers obviate the need for sample dilution and increase sensitivities and dynamic ranges 10-fold. In comparisons against conventional mixing using the same 10 mug of starting total RNA on 22 000-probe arrays, 10 000 more usable signals rose above the noise. In other experiments, planetary mixing was able to

produce comparable results while using only one-tenth the starting sample. The benefits of planetary mixing include sample conservation, shorter hybridizations, less reliance on amplification, and the ability to quantify many gene signals otherwise obscured by noise. 19 Refs.

L3 ANSWER 10 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 2

ACCESSION NUMBER: 2005:1301269 CAPLUS

DOCUMENT NUMBER: 145:226872

TITLE: A microfluidic electrochemical cell based on

microsystem packaging technologies applicable for

biosensor development

AUTHOR(S): Santha, Hunor; Harsanyi, Gabor; Sinkovics, Balint;

Makai, Dora

CORPORATE SOURCE: Department of Electronics Technology, Budapest

University of Technology and Economics, Budapest,

H-1111, Hung.

SOURCE: Proceedings - Electronic Components & Technology

Conference (2005), 55th(Vol. 1), 588-592

CODEN: PETCES

PUBLISHER: Institute of Electrical and Electronics Engineers

DOCUMENT TYPE: Journal LANGUAGE: English

AB A new miniaturized electrochem. cell has been designed and constructed by utilization of different electronics technologies The 25-50 µl capacity of the cylindrical shaped electrochem. cell has been built up by cutting circles in 5-10 overlapping layers of 400 µm thick Teflon foil. This multilayer structure enables very flexible solns. for in- and outlet of different materials into the electrochem. cell. The number of the working electrodes of the biosensor substrates can vary between 1-80. The reference electrode has been made of a silver wire. The circular symmetry of the counter and reference electrodes and the pattern of the working electrode array allows bipotentiostatic measurements, thus, this flexible platform is suitable for wide range of expts. of both bipotentiostatic biocatalytic sensors and DNA sensors with electronically addressed immobilization.

REFERENCE COUNT: 9 · THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 11 OF 35 COMPENDEX COPYRIGHT 2006 EEI on STN

ACCESSION NUMBER: 2005(38):8373 COMPENDEX

TITLE: A microfluidic electrochemical cell based on

microsystem packaging technologies applicable for

biosensor development.

AUTHOR: Santha, Hunor (Department of Electronics Technology

Budapest University of Technology and Economics, Budapest, H - 1111, Hungary); Harsanyi, Gabor;

Sinkovics, Balint; Makai, Dora

MEETING TITLE: 55th Electronic Components and Technology Conference,

ECTC.

MEETING ORGANIZER: IEEE Components, Packaging and Manufacturing

Technology Society; Electronic Components, Assemblies and Materials Association, ECA; Electronic Industries

Alliance, EIA

MEETING LOCATION: Lake Buena Vista, FL, United States

MEETING DATE: 31 May 2005-04 Jun 2005

SOURCE: Proceedings - Electronic Components and Technology

Conference v 1 2005.p 588-592, (IEEE cat n CH37635)

SOURCE: 2005 Proceedings - 55th Electronic Components and

Technology Conference, ECTC

CODEN: PECCA7 ISSN: 0569-5503

PUBLICATION YEAR: 2005 MEETING NUMBER: 65596

DOCUMENT TYPE: Conference Article

TREATMENT CODE: Experimental

LANGUAGE: English
AN 2005(38):8373 COMPENDEX

AB A new miniaturized electrochemical cell has been designed and constructed by utilization of different electronics technologies The 25-50 mul capacity of the cylindrical shaped electrochemical cell has been built up by cutting circles in 5-10 overlapping layers of 400 mum thick Teflon foil. This multilayer structure enables very flexible solutions for inand outlet of different materials into the electrochemical cell. The number of the working electrodes of the biosensor substrates can vary between 1-80. The reference electrode has been made of a silver wire. The circular symmetry of the counter and reference electrodes and the pattern of the working electrode array allows bipotentiostatic measurements, thus, this flexible platform is suitable for wide range of experiments of both bipotentiostatic biocatalytic sensors and DNA sensors with electronically addressed immobilization \$CPY 2005 IEEE. 9 Refs.

L3 . ANSWER 12 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER: 2005:8502840 INSPEC

DOCUMENT NUMBER: A2005-17-6475-019; B2005-09-2575-008

TITLE: Batch-mode mixing on centrifugal microfluidic

platforms

AUTHOR: Grumann, M.; Geipel, A.; Riegger, L.; Zengerle, R.;

Ducree, J. (IMTEK, Freiburg Univ., Germany)

SOURCE: Lab on a Chip (May 2005), vol.5, no.5, p. 560-5, 47

refs.

CODEN: LCAHAM, ISSN: 1473-0197

SICI: 1473-0197 (200505) 5:5L.560:BMMC; 1-6

Published by: R. Soc. Chem, UK

DOCUMENT TYPE: Journal

TREATMENT CODE: Practical; Experimental

COUNTRY: United Kingdom

LANGUAGE: English

DN A2005-17-6475-019; B2005-09-2575-008 AN2005:8502840 INSPEC AB We present two novel fluidic concepts to drastically accelerate the process of mixing in batch-mode (stopped-flow) on centrifugal microfluidic platforms. The core of our simple and robust setup exhibits a microstructured disk with a round mixing chamber rotating on a macroscopic drive unit. In the first approach, magnetic beads which are prefilled into the mixing chamber are periodically deflected by a set of permanent magnets equidistantly aligned at spatially fixed positions in the lab-frame. Their radial positions alternatingly deviate by a slight positive and negative offset from the mean orbit of the chamber to periodically deflect the beads inbound and outbound during rotation. Advection is induced by the relative motion of the beads with respect to the liquid which results from the magnetic and centrifugal forces, as well as inertia. In a second approach - without magnetic beads - the disk is spun upon periodic changes in the sense of rotation. This way, inertia effects induce stirring of the liquids. As a result, both strategies accelerate mixing from about 7 minutes for mere diffusion to less than five seconds. Combining both effects, an ultimate mixing time of less than one second could be achieved

L3 ANSWER 13 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER: 2005:8409489 INSPEC DOCUMENT NUMBER: B2005-06-2575D-026

TITLE: Design and characterization of a passive recycle

micromixer

AUTHOR: Min Ku Jeon; (Dept. of Chem., Korea Adv. Inst. of

Sci. & Technol., Daejeon, South Korea), Joon-Ho Kim; Noh, J.; Soo Ho Kim; Hyun Gyu Park; Seong Ihl Woo

SOURCE: Journal of Micromechanics and Microengineering (Feb.

2005), vol.15, no.2, p. 346-50, 12 refs.

CODEN: JMMIEZ, ISSN: 0960-1317

SICI: 0960-1317 (200502) 15:2L.346:DCPR;1-J

Price: 0960-1317/05/020346+05\$30.00 Doc.No.: S0960-1317(05)83628-9 Published by: IOP Publishing, UK

DOCUMENT TYPE:

Journal

TREATMENT CODE:

Practical; Theoretical

COUNTRY: United Kingdom

LANGUAGE: English

DN B2005-06-2575D-026 AN 2005:8409489 INSPEC

A new design was devised for a recycle micromixer, i.e., a passive AB micromixer with side channels for a recycle flow. The geometry, required to perform a recycle flow and effective mixing, was determined by a simulation based on computational fluid dynamics. A recycle flow of the mixed flow of each unit was introduced to the inlet flow, and a circular flow was generated in the body of the mixer. For complete mixing, five units of the micromixer were connected in series. The simulations were performed at Reynolds numbers of 7, 14 and 28 and channel depths of 100, 150 and 200 µm. Mixing efficiency and direction of recycle flow were significantly affected by both Re and channel depth. When channel depth was 150 µm, mixing efficiency of the micromixer increased from 89.3 to 95.6, 98.4 and 98.6% with the increase of Re from 7 to 14, 28 and 42, respectively. The increasing channel depth also increased mixing efficiency. The micromixer was fabricated by a conventional photolithography technique using polydimethylsiloxane. Color dispersion in blue ink was compared with simulated flow patterns. The characterization of mixing in the recycle micromixer was verified by using an aqueous NaOH solution and phenolphthalein solution, composed of the same volume of ethanol and water. For both cases, fully mixed profiles were achieved along five micromixers, connected in a series at a flow rate of 0.1 ml min-1 for each flow and a short residence time of 0.11 s

ANSWER 14 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER:

2005:8487635 INSPEC

DOCUMENT NUMBER: TITLE:

A2005-16-8780B-040; B2005-08-7230J-060

Common platform for bipotentiostatic biocatalytic

sensors and DNA sensors with electronically addressed

immobilization

AUTHOR:

Santha, H.; Harsanyi, G.; Sinkovics, B.; Takacs, A.

(Dept. of Electron. Technol., Budapest Univ. of

Technol. & Econ., Hungary)

SOURCE:

27th International Spring Seminar on Electronics Technology (IEEE Cat. No.04EX830), vol.1, 2005, p.

136-40 vol.1 of 3 vol. 580 pp., 9 refs.

ISBN: 0 7803 8422 9

Price: 0-7803-8422-9/04/\$20.00

Published by: IEEE, Piscataway, NJ, USA

Conference: 27th International Spring Seminar on Electronics Technology, Bankya, Bulgaria, 13-16 May

DOCUMENT TYPE:

Conference; Conference Article

Practical; Experimental TREATMENT CODE:

United States COUNTRY:

LANGUAGE:

English DN A2005-16-8780B-040; B2005-08-7230J-060 2005:8487635 INSPEC

A new miniaturized electrochemical cell has been designed and constructed by utilization of different electronics technologies. The 25-50 µl capacity of the cylindrical shaped electrochemical cell has been built up by cutting circles in 5-10 overlapping layers of 400 µm thick Teflon foil. This multilayer structure enables very flexible solutions for inlet and outlet of different materials into the electrochemical cell. The number of working electrodes of the biosensor substrates can vary between 1-80. The reference electrode has been made of a silver

wire. The circular symmetry of the counter and reference electrodes and the pattern of the working electrode array allows bipotentiostatic measurements, thus, this flexible platform is suitable for a wide range of experiments of both bipotentiostatic biocatalytic sensors and DNA sensors with electronically addressed immobilization

ANSWER 15 OF 35 INSPEC (C) 2006 IET on STN L3

ACCESSION NUMBER: DOCUMENT NUMBER:

2005:8329022 INSPEC A2005-09-8780-005

TITLE:

A novel high aspect ratio microfluidic

design to provide a stable and uniform

microenvironment for cell growth in a high throughput

mammalian cell culture array

AUTHOR:

Hung, P.J.; Lee, P.J.; Sabounchi, P.; Aghdam, N.; Lin,

R.; Lee, L.P. (Dept. of Bioeng., California Univ.,

Berkeley, CA, USA)

SOURCE:

Lab on a Chip (Jan. 2005), vol.5, no.1, p. 44-8, 19

refs.

CODEN: LCAHAM, ISSN: 1473-0197

SICI: 1473-0197 (200501) 5:1L.44:NHAR;1-T

Published by: R. Soc. Chem, UK

DOCUMENT TYPE:

Journal Practical

TREATMENT CODE: COUNTRY:

United Kingdom

LANGUAGE:

English

AN 2005:8329022 INSPEC

DN A2005-09-8780-005

We present a high aspect ratio microfluidic device for AB

> culturing cells inside an array of microchambers with continuous perfusion of medium. The device was designed to provide a potential tool for cost-effective and automated cell culture. The single unit of the

array consists of a circular microfluidic chamber 40

μm in height surrounded by multiple narrow perfusion channels 2 μm in height. The high aspect ratio (20) between the microchamber and the perfusion channels offers advantages such as localization of the cells inside the microchamber as well as creating a uniform microenvironment for cell growth. Finite element methods were used to simulate flow profile and mass transfer of the device. Human carcinoma (HeLa) cells were cultured inside the device with continuous perfusion of medium at 37 °C and was grown to confluency. The microfluidic cell

culture array could potentially offer an affordable platform for a wide range of applications in high throughput cell-based screening, bioinformatics, synthetic biology, quantitative cell biology, and systems

biology

ANSWER 16 OF 35 COMPENDEX COPYRIGHT 2006 EEI on STN

ACCESSION NUMBER:

2005(5):1368 COMPENDEX

TITLE:

Continuous perfusion microfluidic cell

culture array for high-throughput cell-based assays. **AUTHOR:** Hung, Paul J. (Berkeley Sensor and Actuator Center

Department of Bioengineering University of California, Berkeley, CA 94720, United States); Lee, Philip J.;

Sabounchi, Poorya; Lin, Robert; Lee, Luke P.

SOURCE:

Biotechnology and Bioengineering v 89 n 1 Jan 5 2005

2005.p 1-8

CODEN: BIBIAU ISSN: 0006-3592

PUBLICATION YEAR:

2005 Journal

DOCUMENT TYPE:

Theoretical; Experimental

TREATMENT CODE:

LANGUAGE:

English

2005(5):1368 COMPENDEX ΝA

AB We present for the first time a microfluidic cell culture array for long-term cellular monitoring. The 10 * 10 array could potentially assay 100 different cell-based experiments in parallel. The device was designed to integrate the processes used in typical cell culture experiments on a single self-contained microfluidic system. Major functions include repeated cell growth/passage cycles, reagent introduction, and real-time optical analysis. The single unit of the array consists of a circular microfluidic chamber, multiple narrow perfusion channels surrounding the main chamber, and four ports for fluidic access. Human carcinoma (HeLa) cells were cultured inside the device with continuous perfusion of medium at 37deg C. The observed doubling time was 1.4 +- 0.1 days with a peak cell density of [similar to]2.5*105 cells/cm2. Cell assay was demonstrated by monitoring the fluorescence localization of calcein AM from 1 min to 10 days after reagent introduction. Confluent cell cultures were passaged within the microfluidic chambers using trypsin and successfully regrown, suggesting a stable culture environment suitable for continuous operation. The cell culture array could offer a platform for a wide range of assays with applications in drug screening, bioinformatics, and quantitative cell biology. \$CPY 2004 Wiley Periodicals, Inc. 22 Refs.

L3 ANSWER 17 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 3

ACCESSION NUMBER:

2004:1056511 CAPLUS

DOCUMENT NUMBER:

142:186390

TITLE:

Fabrication of circular-type microchannel using

photoresist reflow and isotropic etching for

microfluidic devices

AUTHOR (S):

Seo, Chang-Taeg; Bae, Chang-Hyun; Eun, Duk-Soo; Shin,

Jang-Kyoo; Lee, Jong-Hyun

CORPORATE SOURCE:

Department of Electronics, Kyungpook National

University, Taegu, 702-701, S. Korea

SOURCE:

Japanese Journal of Applied Physics, Part 1: Regular Papers, Short Notes & Review Papers (2004), 43(11A),

7773-7776 CODEN: JAPNDE

PUBLISHER:

Japan Society of Applied Physics

DOCUMENT TYPE:

Journal

LANGUAGE:

English

AB The authors fabricated circular channels using photoresist reflow and isotropic etching. A silicon substrate, using Si3N4 as a mask, was selectively etched using the hydrofluoric acid, nitric acid, acetic acid (HNA) etching system for fabricating the bottom hemisphere of the channels. Photoresist reflow was used to make the top of the channels round. Then Si3N4 was deposited on the reflowed photoresist. Since the deposited Si3N4 was approx. 6000 Å thick, it was possible to observe the inside of the channel. The authors expect to apply such circular channels to simple bio-systems and microfluidic devices in which optical detection is required.

REFERENCE COUNT:

THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 18 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER: DOCUMENT NUMBER:

2004:8130580 INSPEC A2004-22-8280-026

TITLE:

Dynamic interfacial effect of electroosmotic slip flow with a moving capillary front in hydrophobic circular

microchannels

AUTHOR:

Jun Yang; Fuzhi Lu; Kwok, D.Y. (Dept. of Mech. Eng.,

Univ. of Alberta, Edmonton, Alta., Canada)

SOURCE:

Journal of Chemical Physics (15 Oct. 2004), vol.121,

no.15, p. 7443-8, 53 refs. CODEN: JCPSA6, ISSN: 0021-9606

SICI: 0021-9606(20041015)121:15L.7443:DIEE;1-F Price: 0021-9606/2004/121(15)/7443(6)/\$22.00

Doc.No.: S0021-9606(04)70839-9

Published by: AIP, USA

DOCUMENT TYPE:

Journal

TREATMENT CODE: Theoretical COUNTRY: United States

LANGUAGE: English

AN 2004:8130580 INSPEC DN A2004-22-8280-026

AB Miniaturization of chemical analysis using microfabrication is an emerging technology. The use of polymeric materials as opposed to conventional glass substrate is also a promising alternative. As most polymeric materials are hydrophobic relative to glass, we describe here the implication for the loading process of electroosmotic flow (EOF) when a three-phase (solid-liquid-vapor) contact line exists. The presence of these interfaces can result in a large Laplace pressure that resists EOF and hence hinders its flow performance. This effect depends on the phenomenological contact angle at the solid-liquid interface. In our model for EOF, we considered simultaneously the presence of an electric double layer, liquid slips via a weaker solid-liquid interaction and Laplace pressure across a liquid-vapor interface

L3 ANSWER 19 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2004:874869 CAPLUS

DOCUMENT NUMBER: 142:34613

TITLE: Hybridization Enhancement Using Microfluidic

Planetary Centrifugal Mixing

AUTHOR(S): Bynum, Magdalena A.; Gordon, Gary B.

CORPORATE SOURCE: Agilent Laboratories, Agilent Technologies, Palo Alto,

CA, 94304, USA

SOURCE: Analytical Chemistry (2004), 76(23), 7039-7044

CODEN: ANCHAM; ISSN: 0003-2700

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

DNA microarrays produce their greatest sensitivities when hybridized using AB concentrated samples and effective mixing; however, these goals have proved elusive to combine. If samples are diluted enough to fill larger chambers, then mixing works well using either pumping or gravity with rotation, although sensitivities will suffer. Various techniques for mixing concentrated samples in small thin chambers have been proposed; however, they often leave streaks or scars, and their reusable components require careful cleaning. Here we introduce a versatile new microfluidics platform, a two-axis centrifuge whose fluidic chambers rotate in a planetary relationship to a radial gravitational field. This paradigm readily overcomes surface and viscous forces even in chambers only 50 µm thin. Thin chambers obviate the need for sample dilution and increase sensitivities and dynamic ranges 10-fold. In comparisons against conventional mixing using the same 10 µg of starting total RNA on 22 000-probe arrays, 10 000 more usable signals rose above the noise. In other expts., planetary mixing was able to produce comparable results while using only one-tenth the starting sample. The benefits of planetary mixing include sample conservation, shorter hybridizations, less reliance on amplification, and the ability to quantify many gene signals otherwise obscured by noise.

REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 20 OF 35 COMPENDEX COPYRIGHT 2006 EEI on STN

ACCESSION NUMBER: 2004(30):3478 COMPENDEX

TITLE: A novel magnetic tweezers for manipulation of a single

DNA molecule.

AUTHOR: Chiou, Chi-Han (Department of Engineering Science

National Cheng Kung University, Tainan 701, Taiwan);

Tseng, Zhao-Fu; Lee, Gwo-Bin

MEETING TITLE: 17th IEEE International Conference on Micro Electro

Mechanical Systems (MEMS): Maastricht MEMS 2004

Technical Digest.

MEETING ORGANIZER: IEEE, Robotics and Automation Society

MEETING LOCATION:

Maastricht, Netherlands 25 Jan 2004-29 Jan 2004

MEETING DATE: SOURCE:

Proceedings of the IEEE International Conference on

Micro Electro Mechanical Systems (MEMS) 2004.p

613-616, (IEEE cat n 04CH37517) CODEN: PMEME5 ISSN: 1084-6999

PUBLICATION YEAR: MEETING NUMBER:

2004 63290

DOCUMENT TYPE: Conference Article

TREATMENT CODE: Experimental

LANGUAGE: English AN 2004(30):3478 COMPENDEX

AB We report a novel magnetic tweezers for manipulation of a single DNA molecule. The micromachined DNA manipulator can stretch and rotate a single DNA molecule using arrayed microcoils. Key platform technologies including localized DNA immobilization, microcoil fabrication and microfluidics, have been integrated to form the magnetic DNA tweezers. A single DNA molecule is specifically attached onto a magnetic bead and a gold surface and manipulated under a magnetic field generated by built-in hexagonally-aligned microcoils. A highly effective method for the construction of DNA two sticky ends is developed, which is compatible with MEMS technologies. We have successfully demonstrated the rotation of the tethered-bead DNA molecule linked to the gold pattern by circular permutation of the currents applied to the microcoils. 9

L3 ANSWER 21 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER:

2004:8032541 INSPEC

DOCUMENT NUMBER:

Refs.

A2004-17-8715-027; B2004-08-2575D-087

TITLE:

A novel magnetic tweezers for manipulation of a single

DNA molecule

AUTHOR:

SOURCE:

Chi-Han Chiou; Zhao-Fu Tseng; Gwo-Bin Lee (Dept. of Eng. Sci., Nat. Cheng Kung Univ., Tainan, Taiwan) 17th IEEE International Conference on Micro Electro Mechanical Systems. Maastricht MEMS 2004 Technical Digest (IEEE Cat. No.04CH37517), 2004, p. 613-16 of

li+868 pp., 9 refs. ISBN: 0 7803 8265 X

Price: 0 7803 8265 X/2004/\$17.00

Published by: IEEE, Piscataway, NJ, USA

Conference: 17th IEEE International Conference on Micro Electro Mechanical Systems. Maastricht MEMS 2004 Technical Digest, Maastricht, Netherlands, 25-29 Jan.

2004

Sponsor(s): IEEE; Robotics and Automation Soc

DOCUMENT TYPE: Conference; Conference Article

TREATMENT CODE:

Practical; Experimental

COUNTRY:

United States

LANGUAGE:

English

2004:8032541 INSPEC DN A2004-17-8715-027; B2004-08-2575D-087 AN AB We report a novel magnetic tweezers for manipulation of a single DNA molecule. The micromachined DNA manipulator can stretch and rotate a single DNA molecule using arrayed microcoils. Key platform technologies including localized DNA immobilization, microcoil fabrication and microfluidics, have been integrated to form the magnetic DNA tweezers. A single DNA molecule is specifically attached onto a magnetic bead and a gold surface and manipulated under a magnetic field generated by built-in hexagonally-aligned microcoils. A highly effective method for the construction of DNA two sticky ends is developed, which is compatible with MEMS technologies. We have successfully demonstrated the rotation of the tethered-bead DNA molecule linked to the gold pattern by circular permutation of the currents applied to the microcoils

L3 ANSWER 22 OF 35 COMPENDEX COPYRIGHT 2006 EEI on STN DUPLICATE 4

ACCESSION NUMBER: 2004(50):5073 COMPENDEX

TITLE: NanoLiterBioReactor: Long-term mammalian cell culture

at nanofabricated scale.

AUTHOR: Prokop, Ales (NanoDelivery Inc., Nashville, TN 37211,

United States); Prokop, Zdenka; Schaffer, David; Kozlov, Eugene; Wikswo, John; Cliffel, David;

Baudenbacher, Franz

SOURCE: Biomedical Microdevices v 6 n 4 December 2004 2004.p

325-339

CODEN: BMICFC ISSN: 1387-2176

PUBLICATION YEAR: 2004
DOCUMENT TYPE: Journal
TREATMENT CODE: Experimental
LANGUAGE: English

AN 2004 (50):5073 COMPENDEX

There is a need for microminiaturized cell-culture environments, i.e. AB NanoLiter BioReactors (NBRs), for growing and maintaining populations of up to several hundred cultured mammalian cells in volumes three orders of magnitude smaller than those contained in standard multi-well screening plates. These devices would enable the development of a new class of miniature, automated cell-based bioanalysis arrays for monitoring the immediate environment of multiple cell lines and assessing the effects of drug or toxin exposure. We fabricated NBR prototypes, each of which incorporates a culture chamber, inlet and outlet ports, and connecting microfluidic conduits. The fluidic components were molded in polydimethylsiloxane (PDMS) using soft-lithography techniques, and sealed via plasma activation against a glass slide, which served as the primary culture substrate in the NBR. The input and outlet ports were punched into the PDMS block, and enabled the supply and withdrawal of culture medium into/from the culture chamber (10-100 nL volume), as well as cell seeding. Because of the intrinsically high oxygen permeability of the PDMS material, no additional CO2/air supply was necessary. The developmental process for the NBR typically employed several iterations of the following steps: Conceptual design, mask generation, photolithography, soft lithography, and proof-of-concept culture assay. We have arrived at several intermediate designs. One is termed "circular NBR with a central post (CP-NBR), " another, "perfusion (grid) NBR (PG-NBR), " and a third version, "multitrap (cage) NBR (MT-NBR)," the last two providing total cell retention. Three cells lines were tested in detail: a fibroblast cell line, CHO cells, and hepatocytes. Prior to the culturing trials, extensive biocompatibility tests were performed on all materials to be employed in the NBR design. To delineate the effect of cell seeding density on cell viability and survival, we conducted separate plating experiments using standard culture protocols in well-plate dishes. In both experiments, PicoGreen assays were used to evaluate the extent of cell growth achieved in 1-5 days following the seeding. Low seeding densities resulted in the absence of cell proliferation for some cell lines because of the deficiency of cell-cell and extracellular matrix (ECM)-cell contacts. High viabilities were achieved in all designs. We conclude that an instrumented microfluidics-based NanoBioReactor (NBR) will represent a dramatic departure from the standard culture environment. The employment of NBRs for mammalian cell culture opens a new paradigm of cell biology, so far largely neglected in the literature. \$CPY 2004 Kluwer Academic Publishers. 58 Refs.

L3 ANSWER 23 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER: 2004:7904349 INSPEC DOCUMENT NUMBER: A2004-09-8245-010

TITLE: Design and experiment research of integrated capillary

electrophoresis chip

AUTHOR: Tian Li; Liu Xiao-Wei; Wang Xi-Lian; (MEMS Center,

Harbin Inst. of Technol., China), Tian Lei; Ao

Ming-Sheng; Xiong Jun

SOURCE: Micronanoelectronic Technology (2003), vol.40, no.7-8,

p. 340-3, 4 refs.ISSN: 1671-4776

SICI: 1671-4776(2003)40:7/8L.340:DERI;1-A

Published by: Editorial Board of Micronanoelectronic

Technol, China

DOCUMENT TYPE:

Journal

TREATMENT CODE:

Practical; Experimental

COUNTRY:

China

LANGUAGE:

Chinese

AN 2004:7904349 INSPEC

DN A2004-09-8245-010

AB The fluid distribution in the micro-channel of ICEC (integrated capillary electrophoresis chip) was simulated using ANSYS software. The relations between the micro-channel structure and flow velocity under different sample injection modes were obtained. Then the optimized channel structure was chosen by the width of 16 μm, the depth of the chip was determined 10 μm and the effective separation length of 3.5 cm with round corner channel. The chip structure was designed. Using the principle of laser-induced-fluorescence, the detection system for this chip was set up. The experiment for the ICEC could realize the base line separation of two DNA restriction fragments ranging from 500 bp to 1021 bp with excellent resolution. The result of the experiment would be the stable groundwork for further research in ICEC

L3 ANSWER 24 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER: DOCUMENT NUMBER:

2004:7947668 INSPEC B2004-06-2575F-013

TITLE:

Fabrication of cylinder type micro channel using

photoresist reflow and isotropic etching

AUTHOR:

Chang-Hyun Bae; Chang-Taeg Seo; Jong-Hyun Lee (Sch. of Electron. & Electr. Eng., Kyungpook Nat. Univ., Taegu,

South Korea)

SOURCE:

Digest of Papers Microprocesses and Nanotechnology

2003. 2003 International Microprocesses and

Nanotechnology Conference, 2003, p. 200-1 of xxv+349

pp., 4 refs.

ISBN: 4 89114 040 2

Published by: Japan Soc. of Appl. Phys, Tokyo, Japan

Conference: Digest of Papers Microprocesses and

Nanotechnology 2003. 2003 International Microprocesses and Nanotechnology Conference, Tokyo, Japan, 29-31

Oct. 2003

Sponsor(s): Japan Soc. of Appl. Phys

DOCUMENT TYPE:

Conference; Conference Article

TREATMENT CODE:

Practical; Experimental

COUNTRY:

Japan

LANGUAGE:

English

AN 2004:7947668 INSPEC

DN B2004-06-2575F-013

AB In this paper we have fabricated cylinder type micro channel using photoresist reflow and isotropic etching. Si substrate, using a Si3N4 (6000A) as a mask, was selectively etched using a HNA etching system for the bottom hemisphere of the channel. Photoresist reflow was used to make the top of the channel round. Then Si3N4 was deposited on the reflowed photoresist. Since the deposited Si3N4 was about 6000 A thick it is possible to see the inside of the channel. This channel is expected to be applied to simple bio and microfluidic devices in which optical detection is needed

L3 ANSWER 25 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER:

2004:7937855 INSPEC

DOCUMENT NUMBER:

A2004-11-6180B-007; B2004-05-2575-050 Laser micromachining for microfluidic,

microelectronic and MEMS applications

AUTHOR:

TITLE:

Fedosejevs, R.; Argument, M.; Sardarli, A.; Kirkwood, S.E.; Holenstein, R.; Tsui, Y.Y. (Dept. of Electr. &

Comput. Eng., Univ. of Alberta, Edmonton, Alta,

Canada)

SOURCE: Proceedings International Conference on MEMS, NANO and

Smart Systems, 2003, p. 53 of xii+437 pp.

Editor(s): Badawy, W.; Moussa, W.

ISBN: 0 7695 1947 4

Price: 0 7695 1947 4/2003/\$17.00

Published by: IEEE Comput. Soc, Los Alamitos, CA, USA Conference: Proceedings International Conference on MEMS, NANO and Smart Systems, Banff, Alta., Canada,

20-23 July 2003

Sponsor(s): iCore, Canda; MANCEF, USA; Micronet R&D,

Canada; Nat. Inst. Nano Technol., Canada; Univ.

Alberto, Canada; Univ. Calgary, Canada

Conference; Conference Article

TREATMENT CODE: Application; Practical; Theoretical

COUNTRY: United States

LANGUAGE: English

DOCUMENT TYPE:

AB

AN 2004:7937855 INSPEC DN A2004-11-6180B-007; B2004-05-2575-050

'Summary form only given'. Laser micromachining of dielectrics and metals is a versatile fabrication and repair tool for applications in micro-fluidics, microelectronics and MEMS. Issues such as ablation threshold, ablation rate, incubation of damage at subthreshold fluences, edge resolution, debris creation and residual substrate damage are all important in determining the suitability of this technique for these applications. An extensive study is being carried out on drilling of holes in glasses of interest to microfluidic systems using both nanosecond UV laser pulses at 266 nm and femtosecond pulses at 800 nm. The drill rate and maximum ablation depth has been measured for various holes sizes ranging from 25 to 100 μm in diameter. The resultant morphology is measured using an optical microscope and a scanning electron microscope showing that redeposition and loss of beam fluence down the hole limits the maximum hole depth for a given entrance laser fluence. The results indicate that, in some cases at deep depths near the end point of the hole, the shape may no longer be round . Cracking around the entrance of the hole is also observed with nanosecond drilling and a new technique involving heating of the substrate during the laser interaction is being studied to try to reduce the degree of cracking. Redeposition of glass around the entrance of the drilled hole, which complicates applications where contact must be made to an adjoining surface, is another issue. In order to minimize redeposition, techniques of using a protective sacrificial layer are being investigated. In particular, the use of a protective tungsten thin film which can be stripped off, taking the overcoated debris with the film, is being investigated. In order to understand the microscopic processes occurring within the material and to predict the expected ablation thresholds, ablation rates and residual damage to the material structure, a molecular dynamics simulation code is under development to model the ablation of silicon and the ablation of glasses. Current

L3 ANSWER 26 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2002:906820 CAPLUS

DOCUMENT NUMBER: 137:392245

this paper

TITLE: Method for producing microchannels having circular

cross-sections in glass for microfluidics

INVENTOR(S): Krulevitch, Peter; Hamilton, Julie K.; Ackler, Harold

experimental results and theoretical understanding will be presented in

D.

PATENT ASSIGNEE(S): The Regents of the University of California, USA

SOURCE: U.S. Pat. Appl. Publ., 5 pp.

CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

APPLICATION NO. DATE PATENT NO. KIND DATE -------------------US 2002174686 A1 20021128 US 2001-851231 20010507 PRIORITY APPLN. INFO.: US 2001-851231 20010507

A process for micromachining capillaries was having circular cross sections in glass substrates. Microchannels are isotropically etched into a flat glass substrate, resulting in a semi-circular half-channel (or a rectangle with rounded corners). A 2nd flat glass substrate is then fusion bonded to the 1st substrate, producing sealed microchannels with rounded bottom corners and a flat top surface having sharp corners. The process is completed by annealing at a sufficiently high temperature (.apprx.750°) to allow surface tension forces and diffusional effects to lower the over-all energy of the microchannels by transforming the cross-section to a circular shape. The process can be used to form microchannels with circular cross sections by etching channels into a glass substrate, then anodically bonding to a Si wafer and annealing. The process will work with other materials such as polymers.

L3 ANSWER 27 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER: 2002:7364985 INSPEC DOCUMENT NUMBER: B2002-10-2575F-027

TITLE: Process characterization of fabricating 3D micro

channel systems by laser-micromachining

AUTHOR: Qin, S.J.; Li, W.J. (Dept. of ACAE, Chinese Univ. of Hong Kong, Shatin, China)

SOURCE: Sensors and Actuators A (Physical) (1 April 2002),

vol.A97-98, p. 749-57, 9 refs. CODEN: SAAPEB, ISSN: 0924-4247

SICI: 0924-4247 (20020401) A9798L.749: PCFM; 1-6

Price: 0924-4247/02/\$22.00 Doc.No.: S0924-4247(02)00016-X Published by: Elsevier, Switzerland

Conference: Proceedings of 11th International Conference on Solid State Sensors and Actuators

Transducers '01/Eurosensors XV, Munich, Germany, 10-14

June 2001

DOCUMENT TYPE: Conference; Conference Article; Journal

TREATMENT CODE: Experimental COUNTRY: Switzerland LANGUAGE: English

AN 2002:7364985 INSPEC DN B2002-10-2575F-027

AB A novel process technology was developed to create 3D micro channel systems bounded by solid 3D quartz substrates without damaging the bounding surfaces of the substrate. The process uses a Nd:YAG laser to induce thermal energy or plasma to micromachine channels in substrates which are transparent to the spectrum from UV to near IFR wavelength. We have demonstrated that this process is capable of fabricating up to 4 mm long circular cross-section channels with diameters ranging from 25 to 200 µm. The channel diameter can be controlled by a software program that interfaces with the laser system, thus allowing complete channel systems to be designed on a CAD software and then directly fabricated by the laser system. The process technology, process characterization, and initial test results of the fabricated micro channels are presented in this paper

L3 ANSWER 28 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER: 2002:7348134 INSPEC DOCUMENT NUMBER: B2002-09-2575F-072

TITLE: Fabrication of complex micro channel systems inside

optically-transparent 3D substrates by laser

processing

AUTHOR: Shui-jie Qin; Li, W.J.; (Center for Micro & Nano

Syst., City Univ. of Hong Kong, China), Tao Mei

SOURCE: TRANSDUCERS '01. EUROSENSORS XV. 11th International Conference on Solid-State Sensors and Actuators.

Conference on Solid-State Sensors and Actuators. Digest of Technical Papers, vol.2, 2001, p. 1624-7

vol.2 of 2 vol. 1807 pp., 8 refs.

Editor(s): Obermeier, E. ISBN: 3 540 42150 5

Published by: Springer-Verlag, Berlin, Germany Conference: Proceedings of 11th International Conference on Solid State Sensors and Actuators

Transducers '01/Eurosensors XV, Munich, Germany, 10-14

June 2001

DOCUMENT TYPE: Conference; Conference Article

TREATMENT CODE: Practical; Experimental

COUNTRY: Germany LANGUAGE: English

AN 2002:7348134 INSPEC DN B2002-09-2575F-072

AB A novel process technology was developed to create 3D micro channel systems bounded by solid 3D quartz substrates without damaging the bounding surfaces of the substrate. The process uses a Nd:YAG laser to induce thermal energy or plasma to micromachine channels in substrates which are transparent to Nd:YAG laser wavelength. We have demonstrated that this process is capable of fabricating up 4 mm long circular cross-section channels with diameters ranging from 25 to 200 microns. The channel diameter can be controlled by a software program that interfaces with the laser system, thus allowing

complete channel systems to be designed on a CAD software and then directly fabricated by a laser system. The process technology, process characterization, and initial test results of the fabricated micro channels are presented in this paper

L3 ANSWER 29 OF 35 COMPENDEX COPYRIGHT 2006 EEI on STN

ACCESSION NUMBER: 2000(27):7060 COMPENDEX

TITLE: Normally closed in-channel micro check valve.

AUTHOR: Wang, Xuan-Qi (California Inst of Technology,

Pasadena, CA, USA); Tai, Yu-Chong

MEETING TITLE: 13th Annual International Conference on Micro Electro

Mechanical Systems (MEMS 2000).

MEETING ORGANIZER: IEEE Robotics and Automation Society
MEETING LOCATION: Miyazaki, Jpn

MIYAZAKI, UP

MEETING DATE: 23 Jan 1900-27 Jan 1900

SOURCE: Proceedings of the IEEE Micro Electro Mechanical

Systems (MEMS) 2000.p 68-73

CODEN: PMEME5

PUBLICATION YEAR: 2000
MEETING NUMBER: 56837
DOCUMENT TYPE: Journal
TREATMENT CODE: Theoretical
LANGUAGE: English

AN 2000(27):7060 COMPENDEX

AB We present here the first surface-micromachined, normally closed, in-channel, Parylene check valve. This device is fabricated monolithically on a silicon substrate using a five-layer Parylene process. The operating structure of the check valve is a circular sealing plate on top of a ring-shaped valve seat. The sealing plate is center-anchored on top of a chamber diaphragm that is vacuum-collapsed to the bottom of the chamber in order to achieve a normally closed position. A thin gold layer on the roughened valve seat surface is used to reduce stiction between the sealing plate and the valve seat. We have achieved an in-channel check valve with a cracking (opening) pressure of 20 to approximately 40 kPa under forward bias and no measurable leakage under reverse bias up to 270 kPa. Using this design, this valve performs well in

two-phase microfluidic systems (i.e.microchannel flows containing gas, liquid, or gas/liquid mixture). (Author abstract) 6 Refs.

ANSWER 30 OF 35 INSPEC (C) 2006 IET on STN 1,3

2000:6636002 INSPEC ACCESSION NUMBER:

DOCUMENT NUMBER: B2000-08-8380M-014; C2000-08-3260P-013

A normally closed in-channel micro check valve TITLE: Xuan-Qi Wang; Yu-Chong Tai (Dept. of Electr. Eng., **AUTHOR:**

California Inst. of Technol., Pasadena, CA, USA) Proceedings IEEE Thirteenth Annual International SOURCE: Conference on Micro Electro Mechanical Systems (Cat.

No.00CH36308), 2000, p. 68-73 of xiv+810 pp., 6 refs.

ISBN: 0 7803 5273 4

Price: 0 7803 5273 4/2000/\$10.00

Published by: IEEE, Piscataway, NJ, USA

Conference: Proceedings IEEE Thirteenth Annual

International Conference on Micro Electro Mechanical

Systems, Miyazaki, Japan, 23-27 Jan. 2000

Sponsor(s): IEEE Robotics & Autom. Soc.; Micromachine

Center

Conference; Conference Article DOCUMENT TYPE:

TREATMENT CODE: Practical; Experimental

COUNTRY: United States

LANGUAGE: English

SOURCE:

DN B2000-08-8380M-014; C2000-08-3260P-013 ΑN 2000:6636002 INSPEC

We present here the first surface-micromachined, normally closed, AB in-channel, Parylene check valve. This device is fabricated

monolithically on a silicon substrate using a five-layer Parylene process. The operating structure of the check valve is a circular sealing plate on top of a ring-shaped valve seat. The sealing plate is center-anchored on top of a chamber diaphragm that is vacuum-collapsed to the bottom of the chamber in order to achieve a normally closed position. A thin gold layer on the roughened valve seat surface is used to reduce stiction between the sealing plate and the valve seat. We have achieved an in-channel check valve with a cracking (opening) pressure of 20 40 kPa under forward bias and no measurable leakage under reverse bias up to 270 kPa. Using this design, this valve

performs well in two-phase microfluidic systems (i.e. microchannel flows containing gas, liquid, or gas/liquid mixture)

ANSWER 31 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER: 2001:6894274 INSPEC DOCUMENT NUMBER: B2001-05-2575F-026

TITLE: Deep X-ray exposure system with multistage for 3D

microfabrication

You, H.; Matsuzuka, N.; Yamaji, T.; Tabata, O. (Fac. AUTHOR:

of Sci. & Eng., Ritsumeikan Univ., Shiga, Japan)

MHS2000. Proceedings of 2000 International Symposium

on Micromechatronics and Human Science (Cat. No.00TH8530), 2000, p. 53-8 of vii+247 pp., 7 refs.

ISBN: 0 7803 6498 8 Price: 0 7803 6498 8/2000/\$10.00

Published by: IEEE, Piscataway, NJ, USA

Conference: MHS2000. Proceedings of 2000 International

Symposium on Micromechatronics and Human Science,

Nagoya, Japan, 22-25 Oct. 2000

Sponsor(s): IEEE Ind. Electron. Soc.; IEEE Robotics &

Autom. Soc.; City of Nagoya; Nagoya Urban Ind.

Promotion Corp.; Chubu Ind. Adv. Center; Nagoya Univ.; Chubu Sci. & Technol. Center; Japan Soc. Mech. Eng.; Robotics Soc. Japan; Soc. Instrum. & Control Eng.;

Res. Committee on Micromechatronics; Tech. Committee on Micro-mechanisms of Japan Soc. of Japan Soc.

Precision Eng.; Chubu Bureau of Int. Trade & Ind.

MITI; Federation of Micromachine Technol.; Micromachine Center; Aichi Prefecture; Gifu

Prefecture; Shizuoka Prefecture; Nagano Prefecture;

Nagoya Chamber of Commerce & Ind.; Chubu Econ.

Federation; Nagoya Junior Chamber Conference; Conference Article

TREATMENT CODE: Practical; Experimental

COUNTRY: United States

LANGUAGE: English

DOCUMENT TYPE:

AN 2001:6894274 INSPEC DN B2001-05-2575F-026

AB This paper reports a new deep X-ray exposure system to realize 3D microstructures with controllable curved and inclined walls. Based

microstructures with controllable curved and inclined walls. Based on a compact synchrotron light source, a dedicated X-ray beamline and the exposure device have been constructed. They could work in exposure environments of vacuum or helium gas. The exposure device was mainly made up of 5 stages and had as many as 6 degrees of freedom, which enabled the system to have more functions than the normal one. Besides the scan in its plane, the substrate surface could also rotate round one of its normal and tangent respectively. Driven by a PZT stage, the X-ray mask could move freely in its plan against the substrate behind, which were used to control the wall inclination and flexure of the substrate structure. The system also had off-line mask-substrate alignment function. Various 3D PMMA microstructures can be realized by the system, such as lens array, nozzles, tube connector, conical tubes, inclined channels, long circle channels, angle pipe with smooth joint, cone, gear rack, long column with curve cross-section etc., which are impossible with the normal X-ray lithography system. A series of deep X-ray lithography experiments have been completed and obtained some interesting 3D PMMA microstructures and the relationship between the etching depth and the dose energy of the exposure. It demonstrated the potential of the system, which will greatly enlarge the application fields of deep X-ray lithography and LIGA process

L3 ANSWER 32 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER:

DOCUMENT NUMBER: A20

TITLE:

AUTHOR:

SOURCE:

2000:6515952 INSPEC

A2000-07-0710C-018; B2000-04-2575F-039
Mass separation using thin PTFE membranes

Rummler, Z.; Bacher, W.; Saile, V.; Schomburg, W.K.

(Forschungszentrum, Inst. fur Mikrostrukturtech.,

Karlsruhe, Germany)

Proceedings of the SPIE - The International Society for Optical Engineering (1999), vol.3680, pt.1-2, p.

1014-21, 15 refs.

United States

CODEN: PSISDG, ISSN: 0277-786X

SICI: 0277-786X(1999)3680:1/2L.1014:MSUT;1-#

Price: 0277-786X/99/\$10.00

Published by: SPIE-Int. Soc. Opt. Eng, USA

Conference: Design, Test, and Microfabrication of MEMS

and MOEMS, Paris, France, 30 March-1 April 1999

Sponsor(s): SPIE; CNRS-INPC-UJF; IEEE
Conference; Conference Article; Journal

TREATMENT CODE: Practical; Experimental

COUNTRY:

DOCUMENT TYPE:

LANGUAGE: English

AN 2000:6515952 INSPEC DN A2000-07-0710C-018; B2000-04-2575F-039 Devices for mass separation have been engineered and were fabricated using the AMANDA process. The key component is a 5 μ m thin, semi-permeable PTFE (polytetrafluorethylene) membrane with a circular diameter of 42 mm. The membrane is encapsulated in a PTFE and PEEK (polyetheretherketone) housing. In experiments, this novel device separated a gas flow of approximately 1 μ l/min from a methanol feed stream of 1 ml/min at a pressure difference of 900 hPa. The separation process was simulated in FE-calculations exploiting

analogies between diffusion theory and heat transfer. Mechanical

stability and creeping of the PTFE membrane were investigated as well. All parts in contact with the fluids to be separated are made of chemically inert polymers. As a consequence, a welding process had to be developed for bonding the PTFE membrane to the PTFE housing. This was accomplished with an intermediate FEP (polytetrafluorethylene/hexafluorpropylene) layer. Extension of this bonding technique to other AMANDA products will facilitate fabrication of chemically inert micropumps, -valves, and -sensors

L3 ANSWER 33 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER: 2000:6588707 INSPEC

DOCUMENT NUMBER: B2000-06-8380M-007; C2000-06-3260P-006

TITLE: Microvalve analysis: wall shear and diffuser effects

AUTHOR: Carmona, M.; Marco, S.; Samitier, J.; (CSIC,

Barcelona Univ., Spain), Acero, M.C.; Plaza, J.A.;

Esteve, J.

SOURCE: 1999 International Conference on Modeling and

Simulation of Microsystems, 1999, p. 554-7 of

xviii+690 pp., 11 refs.

ISBN: 0 9666135 4 6

Published by: Computational Publications, Cambridge,

MA, USA

Conference: Proceedings of International Conference on

Modelling and Simulation of Microsystems,

Semiconductors, Sensors and Actuators, San Juan,

Puerto Rico, 19-21 April 1999

Sponsor(s): Integrated Syst. Eng.; IntelliSense Corp.;

MEMSCAP S.A.; Mentor Graphics Corp.; Microcosm

Technol. Inc.; Molecular Simulations Inc

DOCUMENT TYPE: Conference; Conference Article

TREATMENT CODE: Practical; Theoretical

COUNTRY: United States

validate the proposed model

LANGUAGE: English

AN 2000:6588707 INSPEC DN B2000-06-8380M-007; C2000-06-3260P-006

This work deals with the analysis of silicon passive microvalves. The analysed microvalves consist of a circular central mass suspended by two flexible beams anchored on a pyrex substrate.

The mechanical and fluidic analysis are first done by FEM (FLOTRAN) simulations, and afterwards the results are compared to an iterative analytical model, showing good agreement at low flow rates. In this model, we have taken into account viscous losses as well as inertial ones, usually neglected in model extraction and validation. A good fitting is achieved at high flow rates for a diminished inertial losses contribution. Measurements with air have been carried out in order to

L3 ANSWER 34 OF 35 COMPENDEX COPYRIGHT 2006 EEI on STN

ACCESSION NUMBER: 2000(16):3537 COMPENDEX

TITLE: Micromachined flow handling components - micropumps.

AUTHOR: Soerensen, Olaf (Inst fuer Microtechnik Mainz GmbH,

Mainz, Ger); Drese, Klaus S.; Ehrfeld, Wolfgang;

Harman Hans Tookin

Hartmann, Hans-Joachim

MEETING TITLE: Proceedings of the 1999 Chemical Microsensors and

Applications II.

MEETING ORGANIZER:

SPIE

MEETING LOCATION:

Boston, MA, USA

MEETING DATE:

19 Sep 1999-20 Sep 1999

SOURCE:

Proceedings of SPIE - The International Society for

Optical Engineering v 3857 1999.p 52-60

CODEN: PSISDG ISSN: 0277-786X

PUBLICATION YEAR:

1999

MEETING NUMBER: DOCUMENT TYPE: 56253 Journal

TREATMENT CODE:

Experimental

LANGUAGE: English

AN2000(16):3537 COMPENDEX

Microfluidic components, especially micropumps, are essential AB for miniaturized dosing systems, Micro Total Analysis Systems (mu -TAS) and miniaturized labs for medical diagnosis. A variety of micropumps for these different applications have been developed at the Institut fuer Mikrotechnik Mainz GmbH (IMM). Self-filling membrane pumps have been realized in various polymer materials. They consist of two injection-molded parts and a valve membrane with a thickness of only 2 mu m. After structuring the membrane with an excimer laser, a Nd: YAG laser is used to weld the parts. The micropump is driven by a piezoelectric actuator which renders frequencies up to 100 Hz. The maximum flow is 400 mu 1/min water at up to 2.0*105 Pa. The total size of the micropump is 12*12*3 mm3. These measures allow it to be integrated e.g. into dosing-systems for active substances in medical applications. For fluids with high viscosity like lubricants for high performance bearings micro gear pumps are very well suited since their maximum flow rate and backpressure is higher than that of membrane pumps. Consequently, various types of gear pumps internal and external geared systems - have been developed at IMM. The gearwheels have been manufactured by LIGA technology and are of circular or oval shape with typical dimensions of 500 to 1,000 mu m.A micro motor with an outer diameter of only 1.9 mm drives one of the gearwheels. Due to its small size the motor can be integrated into the housing of the pump, leading to micropumps, whose overall size is comparable to the size of a sugar cube. (Author abstract) 16 Refs.

ANSWER 35 OF 35 INSPEC (C) 2006 IET on STN

ACCESSION NUMBER:

DOCUMENT NUMBER:

TITLE:

AUTHOR:

SOURCE:

1999:6375724 INSPEC

B1999-11-8380M-019; C1999-11-3260P-019

A PZT-driven micropump

Xiaohao Zang; Zhaoying Zhou; Xiongying Ye; Yong Li;

Wendong Zhang (Dept. of Precision Instrum. & Mech.,

Tsinghua Univ., Beijing, China)

MHA'98. Proceedings of the 1998 International Symposium on Micromechatronics and Human Science. -Creation of New Industry - (Cat. No.98TH8388), 1998,

p. 269-72 of vii+277 pp., 9 refs.

ISBN: 0 7803 5130 4

Price: 0 7803 5130 4/98/\$10.00

Published by: IEEE, Piscataway, NJ, USA Conference: MHS'98. Proceedings of the 1998

International Symposium on Micromechatronics and Human

Science, Nagoya, Japan, 25-28 Nov. 1998

Sponsor(s): IEEE Ind. Electron. Soc.; IEEE Robotics &

Autom. Soc.; City of Nagoya; Nagoya Urban Ind.

Promotion Corp.; Chubu Ind. Adv. Center; Nagoya Univ.;

Chuba Sci. & Technol. Center; Japan Soc. Mech. Eng.; Robotics Soc. Japan; Soc. Instrum. & Control Eng.; Res. Committee on Micromechatronics; Tech. Committee

on Micro-mechanisms of Japan Soc. Precision Eng.; Chubu Bureau of Int. Trade & Ind. MITI; Federation of Micromachine Technol.; Micromachine Center; Aichi

Prefecture; Gifu Prefecture; Mie Prefecture; Shizuoka Prefecture; Nagano Prefecture; Nagoya Chamber of

Commerce & Ind.; Chubu Econ. Federation; Nagoya Junior

Chamber

DOCUMENT TYPE:

TREATMENT CODE:

COUNTRY: LANGUAGE: Conference; Conference Article

Application; Practical; Experimental

United States

English

1999:6375724 INSPEC DN B1999-11-8380M-019; C1999-11-3260P-019 AN

The design, fabrication and characterization of a PZT-driven micropump is AB presented in this paper. It consists of a chamber, a membrane, two microvalves and a driving mechanism. The thickness of the micropump

membrane is 11 μm . The micropump chamber is round with diameter of 5 mm and depth of 0.4 mm. The microvalves made of single crystallite silicon are used on this micropump as flow direction control elements. The dimension of the valve cover is 1.5 mm+1.0 mm+7.4 μ m and dimension of the valve opening is 200 $\mu m + 200 \ \mu m$. Its capability of flow is more than 10 ml/min at a pressure level of about 10 kPa. The open pressure in the obverse direction is less than 200 Pa while the leakage of reverse direction is almost zero. The features of the microvalve fit the requirement of the micropump well. When the micropump chamber chip and two valve chips are assembled together as a micropump, this micropump is mounted on a metal base and then the PZT bimorph cantilever is mounted above the micropump membrane. The maximum flow rate of the micropump is 365 μl/min under 100 V, 20 Hz square wave power supply and zero pressure fall. The back pressure is 2.38 kPa and the flow control precision is better than 1 µl